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(54) Title: NON-AQUEOUS SUSPENSION CONCENTRATE

## (57) Abstract

The invention relates to a non-aqueous, stable suspension concentrate which comprises (a) 50 to 400 g/L of one or more crop protection active compounds, (b) 50 to 700 g/L of one or more adjuvants, (c) 75 to 500 g/L of one or more organic solvents, at least one dispersant selected from the groups (d) and (e), (d) 5 to 150 g/L of one or more non-ionic dispersants, (e) up to 150 g/L of one or more anionic dispersants, and optionally (f) up to 100 g/L of one or more thickeners, and to the pesticidal use of such a suspension.

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## **NON-AQUEOUS SUSPENSION CONCENTRATE**

### **BACKGROUND OF THE INVENTION**

This invention concerns a non-aqueous, stable suspension concentrate (SC) for crop protection active compounds, a method for the manufacture of such suspensions, and their use for combating pests.

As a rule, inert ingredients must be used to provide crop protection active compounds, for example fungicidal compounds, in a form that the user can apply, either as such or after dilution with water. The right choice of suitable inert ingredients, such as carriers, for the formulation often determines to a significant extent whether the active ingredient exhibits its full efficacy on application. Not every active ingredient is suitable for use in any given formulation, because both the efficacy and physiochemical stability of the active ingredient may be affected by other ingredients in the formulation.

The efficacy of the active components can often be improved by addition of other ingredients. The observed efficacy of the combination of ingredients can sometimes be significantly higher than that would be expected from the amounts of the individual ingredients used (synergism). An adjuvant is defined here as a substance which can increase the biological activity of an active ingredient but is not itself significantly biologically active. The adjuvant can either be included in the formulation or can be added separately, e.g., to a spray tank together with the formulation containing the active ingredient.

For easy and safe handling and dosing of these adjuvants by the end-user, and to avoid unnecessary packing material, it is desirable to develop concentrated formulations which already contain such adjuvants.

The International Patent Application WO 95/01722 discloses aqueous pesticidal formulations containing non-ionic dispersants selected

from block-polymers having a polyoxypropylene core, an anionic dispersant, and a wetting agents selected from polyalkoxylated fatty alcohols. However, it is not stated that these wetting agents enhance the activity of the pesticides. Moreover, there is no disclosure of non-aqueous suspension concentrates.

5

### SUMMARY OF THE INVENTION

The present invention relates to a non-aqueous, stable suspension concentrate (SC) which comprises

- 10 (a) 50 to 400 g/L of one or more crop protection active compounds;
- (b) 50 to 700 g/L of one or more adjuvants;
- (c) 75 to 500 g/L of one or more organic solvents;
- at least one dispersant selected from the groups (d) and (e)
- (d) 5 to 150 g/L of one or more non-ionic dispersants,
- 15 (e) 5 to 150 g/L of one or more anionic dispersants; and optionally

it has also surprisingly been found that the biological activity of the active ingredients (a) can be increased by including the adjuvants (b) in the spray dilution or directly in the formulation. In the formulation of the present invention, the adjuvants (b) are incorporated into the concentrated formulation.

5       The term pests as used herein includes, but is not limited to, plant pathogens, insects and weeds.

In principle, all crop protection active compounds can be used in non-aqueous concentrated suspensions according to the invention. Solid crop protection active compounds are preferred.

10       Preferably, the solid crop protection active compounds are less than moderately soluble in the organic solvent (c). A solubility of less than 10 g/L, in particular less than 5 g/L, in solvent (c) is preferred.

The compositions of this invention can be applied to the plants or their environment with additional active substances, such as fertilizers, or agents  
15       containing trace elements, or other preparations which influence plant growth, or

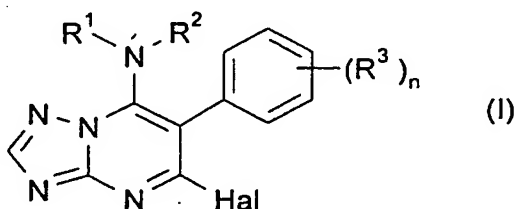
Preferred fungicides for use in the compositions of the present invention are the commercially available compounds selected from the group consisting of:

anilazine, azoxystrobin, benalaxyl, benomyl, binapacryl, bitertanol, blasticidin S, Bordeaux mixture, bromuconazole, bupirimate, captafol, captan, 5 carbendazim, carboxin, carpropamid, chlorbenzthiazon, chlorothalonil, chlozolate, copper-containing compounds such as copper oxychloride, and copper sulfate, cycloheximide, cymoxanil, cypofuram, cyproconazole, cyprodinil, dichlofluanid, dichlone, dichloran, diclobutrazol, diclocymet, diclomezine, diethofencarb, difenoconazole, diflumetorim, dimethirimol, dimethomorph, 10 diniconazole, dinocap, ditalimfos, dithianon, dodemorph, dodine, edifenphos, epoxiconazole, etaconazole, ethirimol, etridiazole, famoxadone, fenapanil, fenamidone, fenarimol, fenbuconazole, fenfuram, fenhexamid, fencpiclonil, fenpropidin, fenpropimorph, fentin, fentin acetate, fentin hydroxide, ferimzone, fluazinam, fludioxonil, flumetover, fluquinconazole, flusilazole, flusulfamide, 15 flutolanil, flutriafol, folpet, fosetyl-aluminium, fuberidazole, furalaxyl, furametpyr,

control of insects, weeds or plant diseases, or to induce host resistance in the plants. Examples of such biological control agents are: *Bacillus thuringiensis*, *Verticillium lecanii*, *Autographica californica* NPV, *Beauveria bassiana*, *Ampelomyces quisqualis*, *Bacillus subtilis*, *Pseudomonas fluorescens*,  
5 *Streptomyces griseoviridis* and *Trichoderma harzianum*.

Moreover, the formulations according to the invention may contain at least one chemical agent that induces the systemic acquired resistance in plants such as, for example, nicotinic acid or derivatives thereof, 2,2-dichloro-3,3-dimethylcylopropylcarboxylic acid, or BION.

10 Also preferred are compositions that include derivatives of triazolopyrimidines which are disclosed, for example, by European Patent Applications EP 0 071 792 and EP-A-0 550 113, in particular the compounds of formula I



15 in which

R<sup>1</sup> and R<sup>2</sup> each independently represent hydrogen or an optionally substituted alkyl, alkenyl, alkynyl, alkadienyl, haloalkyl, aryl, heteroaryl, cycloalkyl, bicycloalkyl or heterocyclyl group, or

R<sup>1</sup> and R<sup>2</sup> together with the adjacent nitrogen atom represent an optionally  
20 substituted heterocyclic ring,

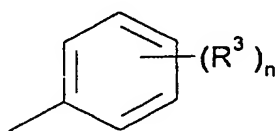
R<sup>3</sup> represents a halogen atom or an alkyl or alkoxy group,

n represents an integer from 0 to 5, and

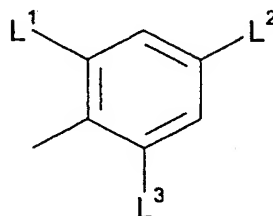
Hal represents a halogen atom.

More preferred are those compounds of formula I wherein R<sup>1</sup> and R<sup>2</sup>  
25 together with the interjacent nitrogen atom represent an optionally substituted 6-membered heterocyclic ring, in particular a 4-methylpiperid-1-yl group; or R<sup>1</sup> represents a C<sub>1-6</sub> alkyl, a C<sub>1-6</sub> haloalkyl, in particular a

$C_{2-6}$  fluoroalkyl, or a  $C_{3-8}$  cycloalkyl group and  $R^2$  represents a hydrogen atom or a  $C_{1-6}$  alkyl group and/or wherein



represents



in which  $L^1$  represents a halogen atom, preferably fluorine or chlorine and  $L^2$  and  $L^3$  each independently represent a hydrogen atom or a halogen atom, preferably fluorine; and/or wherein

Hal represents a chlorine atom.

Most preferred are those compounds of formula I wherein

$R^1$  represents a  $C_{2-5}$  fluoroalkyl group, in particular a 2,2,2-trifluoroethyl or 1,1,1-trifluoroprop-2-yl group; and  $R^2$  represents a hydrogen atom.

Another group of preferred fungicidal compounds are the benzoylbenzenes which are disclosed, for example, by European Patent Application EP-A-0 727 141.

Preferred herbicides include the commercially available compounds selected from the group consisting of:

2,4-D, 2,4-DB, 2,4-DP, acetochlor, acifluorfen, alachlor, alloxydim, ametrydione, amidosulfuron, asulam, atrazin, azimsulfuron, benfuresate, bensulfuron, bentazon, bifenox, bromobutide, bromoxynil, butachlor, cafenstrole, carfentrazone, chloridazon, chlorimuron, chlorpropham, chlorsulfuron, chlortoluron, cinmethylin, cinosulfuron, clomazone, clopyralid, cyanazin, cycloate, cyclosulfamuron, cycloxydim, daimuron, desmedipham, di-methazone, dicamba, dichlobenil, diclofop, diflufenican, dimethenamid, dithiopyr, diuron, eptame, esprocarb, ethiozin, fenoxaprop, flamprop-M-isopropyl, flamprop-M-methyl, fluazifop, fluometuron, fluoroglycofen, fluridone, fluroxypyr, flurtamone, fluthiamid, fomesafen, glufosinate, glyphosate, halosafen, haloxyfop, hexazinone, imazamethabenz, imazamethapyr, imazamox, imazapyr, imazaquin, imazethapyr, ioxynil, isoproturon, isoxaben, isoxaflutole, lactofen, MCPA, MCPP, mefenacet, metabenzthiazuron, metamitron, metazachlor, methyldimron, metolachlor.



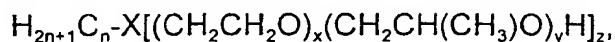
metribuzin, metsulfuron, molinate, nicosulfuron, norflurazon, oryzalin, oxadiargyl, oxasulfuron, oxyfluorfen, pendimethalin, picloram, pretilachlor, propachlor, propanil, prosulfocarb, pyrazosulfuron, pyridate, qinmerac, quinchlorac, quizalofopethyl, sethoxydim, simetryne, sulcotrione, sulfentrazone, sulfosate, terbutryne, terbutylazin, thiameturon, thifensulfuron, thiobencarb, tralkoxydim, triallate, triasulfuron, tribenuron, triclopyr, trifluralin.

Furthermore preferred are the derivatives of aryloxypicolineamides which are disclosed, for example, by European Patent Application EP-A-0 447 004, in particular, N-(4-fluorophenyl) 6-(3-trifluoromethylphenoxy)-pyrid-2-ylcarboxamide having the proposed common name picolinafen.

Examples of insecticidal compounds are alpha-cypermethrin, benfuracarb, BPMC, buprofezine, carbosulfan, cartap, chlorfenvinphos, chlorpyrifos-methyl, cycloprothrin, cypermethrin, esfenvalerate, ethofenprox, fenpropathrin, flucythrinate, flufenoxuron, hydramethylnon, imidacloprid, isoxathion, MEP, MPP, nitenpyram, PAP, permethrin, propaphos, pymetrozine, silafluofen, tebufenozide, teflubenzuron, temephos, terbufos, tetrachlorvinphos and triazamate.

The non-aqueous SC according to the invention comprises 50 to 400 g/L, preferably 75 to 250 g/L, more preferably 80 to 200 g/L of one or more crop protection active compounds.

The adjuvants (b) are preferably liquid polyalkoxylated aliphatic alcohols or amines. These adjuvants may be obtained by alkoxylation of fatty alcohols or amines having 9 - 24, preferably 12 - 22 and in particular 14 - 20 C-atoms, with alkyleneoxide having 2 - 6, preferably 2 - 3 C-atoms, in particular with a mixture of ethylenoxide and propyleneoxide. The aliphatic moieties of the said fatty alcohols and amines may be straight-chained or branched. Preferably these compounds correspond to mixed random or block oligomers of the following formula



in which

X represents O or N,

z is 1, in the event that X represents O, or 2 in the event that X represents N, and the average of the indexes given is as follows:

n is an integer from 9 to 20, in particular 15 to 19;

x is an integer from 1 to 8, in particular 2 to 6; and  
y is an integer from 6 to 12, in particular 7 to 10.

Particularly preferred in the practice of this invention are those polyalkoxylated aliphatic alcohols or amines which are liquids at temperatures down to at least 20°C and have a viscosity of 30 to 100, in particular 50 to 80 mPa · s at 25 °C. The compounds which are commercially available under the trademark Plurafac® LF (Tensid-Chemie, Köln / BASF AG, Ludwigshafen) and certain ATPLUS®-types (ICI Surfactants, Eversberg), in particular Plurafac® LF 224, Plurafac® LF 403, Plurafac® LF 700 and Plurafac® LF 1300, ATPLUS® 245 or SCS4774 (ICI Surfactants) have been proven to be especially advantageous.

In another preferred embodiment of the present invention, the adjuvant (b) is preferably a polyoxyalkylene triglyceride. These adjuvants are obtainable by alkoxylation of triglycerides, resulting in mixtures of compounds with one to three glyceride side chains having 9 - 24, preferably 12 - 22 and in particular 14 - 20 C-atoms, in particular with ethyleneoxide. The aliphatic moieties of the triglycerides may be straight-chained or branched. Preferably, these compounds are mixed oligomers resulting from the alkoxylation of castor or canola oil.

A particularly preferred adjuvant (b) is castor oil ethoxylate, for example Ukanil® 2507, which is commercially available from ICI Surfactants, or canola oil alkoxyate, for example EMULGIN CO3522, which is commercially available from Henkel KGaA.

The non-aqueous SC of this invention comprises 50 to 700 g/L, preferably 200 to 600 g/L, more preferably 300 to 500 g/L of one or more adjuvants.

In a particularly preferred embodiment of the present invention, the adjuvant (b) comprises two or more different alkoxyated derivatives, one of which is an alkoxyated triglyceride, in particular an ethoxylated triglyceride. The non-aqueous SC of this invention preferably comprises 5 to 150 g/L, more preferably 20 to 100 g/L, in particular 40 to 75 g/L of one or more alkoxyated triglyceride, and 45 to 550 g/L of one or more alkoxyated alcohol or amine.

The efficacy of the fungicidal triazolopyrimidines of formula I can be enhanced by addition of adjuvants (b).

In a particularly preferred embodiment, the triazolopyrimidine of formula I is 5-chloro-6-(2-chloro-6-fluorophenyl)-7-(2,2,2-trifluoroethylamino)-[1,2,4]triazolo[1,5-a]pyrimidine (Compound IA), or 5-chloro-6-(2,4,6-trifluorophenyl)-7-(1,1,1-trifluoroprop-1-ylamino)-[1,2,4]triazolo[1,5-a]pyrimidine (Compound IB).

The appropriate relative amounts of active ingredient (a) and the adjuvant (b) lie, in accordance with the invention, between 1:1 and 1:100, preferably between 1:1 and 1:10 and, in particular, between 1:2 and 1:5. Generally, the pesticidal efficacy can be enhanced to a higher degree by the addition of larger amounts of the adjuvant (b), as is shown in the experimental results described below.

Recommended doses for various applications are known for the plant protection active compounds (a); however, the efficacy can be enhanced in accordance with the invention. Addition of the adjuvants of this invention can (depending on the active ingredient, the adjuvant and their amounts) reduce the

1095 (Witco Corp.), preferred alkyl lactates are ethyl and 2-ethylhexyl lactate, preferred glycols are dialkyl diethyleneglycols, in particular diethyl diethyleneglycol. Mixtures of different solvents are often suitable.

The non-aqueous SC according to the invention comprises 75 to 500 g/L, preferably 100 to 450 g/L, in particular 200 to 420 g/L of one or more organic solvents.

The non-ionic dispersant (d) is preferably an ethoxylated non-ionic dispersant different from the adjuvants (b), more preferably polyethyleneoxide-polypropyleneoxide block-copolymers, for example PLURONIC®-type block-copolymers, which are available from BASF AG, or polyoxyethylene fatty acid or polyoxyethylene alcohols. These dispersants are obtainable by alkoxylation of fatty acids, alcohols or alkylphenols having 9 - 24, preferably 12 - 22 and in particular 14 - 20 C-atoms, with ethyleneoxide. The aliphatic moieties of the fatty alcohols may be straight-chained or branched. Preferred dispersants (d) include ARKOPAL®-type alkyl- and/or alkylarylethoxylates (Clariant GmbH former

The non-aqueous SC according to the invention preferably comprises 5 to 150 g/L, more preferably 20 to 100 g/L, in particular 30 to 70 g/L of one or more anionic dispersants.

The thickener (f) is preferably an organo clay or a hydrated silicate, especially a hydrated aluminium magnesium silicate such as Attagel® 50 (Engelhard Corp.) or bentonite derivatives such as BENTONE® SD-1 or SD-3 (Rheox, Inc. Hightstown, NJ, USA), pyrogenic silicic acid such as Car-O-Sil M5 (Cabot GmbH, Rheinfelden, Germany), polyamides or polesters such as THIXATROL® Plus or THIXATROL® 289 (Rheox, Inc. Hightstown, NJ, USA).

The non-aqueous SC according to the invention may comprise up to 100 g/L, preferably 10 to 100 g/L, in particular 30 to 75 g/L of one or more thickeners.

In a particularly preferred embodiment according to this invention the non-aqueous SC essentially consists of

- (a) 75 to 250 g/L of one or more crop protection active compounds, in particular a compound of formula I;
- (b) 200 to 650 g/L of one or more ingredients selected from the group consisting of  
C<sub>9-20</sub> alcohols or amines being alkoxyated with 2 to 20 C<sub>2-6</sub> alkoxy groups, in particular PLURAFAC® LF 700 or ATPLUS® 245, and alkoxyated triglycerides, in particular UKANIL® 2507;
- (c) 100 to 450 g/L of one or more organic solvents selected from the group consisting of hydrocarbons, aliphatic hydrocarbons, alkyl lactates, glycols and plant oil esters;
- (d) 0 to 50 g/L of a polyoxyethylene fatty acid;
- (e) 20 to 100 g/L of an alkali or alkaline earth metal sulfonate, in particular Rhodocal® 70/B or PHENYLSULFONAT CA100; and
- (f) 10 to 100 g/L of one or more silicates or organo clays, in particular Attagel® 50.

The ingredients may be processed to form a suspension concentrate according to the invention by well-established techniques, including intensive mixing and/or milling of the active ingredients with the other substances, such as solvents, dispersants, and adjuvants. The desired form of application, such as

spraying, atomizing, dispersing or pouring, will depend on the desired objectives and the given circumstances, and can be readily determined by one skilled in the art.

Suspension concentrates according to the present invention are usually  
5 produced so as to obtain a stable, non-sedimenting, flowable product containing 5 to 40% w/v active ingredient, 0.5 to 30% w/v of dispersing agents, 0.1 to 10% w/v of suspending agents such as protective colloids and thixotropic agents, 0 to 10% w/v of other additives such as defoamers, corrosion inhibitors, stabilizers, penetrants and stickers, and an organic liquid in which the active ingredient is  
10 substantially insoluble.

In a preferred embodiment, the crop protection compound (a) is air-milled before admixing the components (b) to (f).

The finished non-aqueous suspension concentrates according to the invention are stable in storage, even over a relatively long period. Although phase  
15 separation may occur upon storage due to sedimentation of the active ingredient, no aggregates are formed. The SCs according to the present invention may include high loadings of one or more adjuvants in a one-pack formulation with a pesticide and, therefore, offer the advantage of an optimized and easy-to-use formulation of the crop protection active compound. The separate addition of an  
20 adjuvant by the end-user before application is, therefore, unnecessary.

Aqueous dispersions and emulsions, for example compositions obtained by diluting the SC of the invention with water, also lie within the scope of the invention.

As a commodity, the compositions preferably may be in a concentrated  
25 form, whereas the end user generally employs diluted compositions. The compositions may be diluted to a concentration down to 0.001% of active ingredient. The desired dose usually is in the range from 0.01 to 10 kg a.i./ha.

For a more clear understanding of the invention, specific examples are set forth below. These examples are merely illustrations and are not to be understood  
30 as limiting the scope and underlying principles of the invention in any way. Various modifications of the invention in addition to those shown and described herein will become apparent to those skilled in the art from the following examples

and foregoing description. Such modifications are also intended to fall within the scope of the appended claims.

Examples of non-aqueous suspension concentrates according to the invention are shown in the following examples A to M:

5

Identity of Ingredients used in Examples

Name	Identity
<u>Compound IB</u>	Fungicidal Triazolopyrimidine of formula I
Rhodocal® 70/B (Rhodia)	70% Calcium Dodecylbenzene sulfonate in n-butanol
Ukanil® 2507 (ICI Surfactants)	Castor oil ethoxylate
Emulgin® CO3522 (Henkel)	Canola oil alcoxylate
Plurafac® LF700 (BASF AG)	alcohol alcoxylate
SCS4774 (ICI Surfactants)	branched alcohol alcoxylate
Witconol® 1095 (Witco Corp.)	95% C <sub>10</sub> methylated plant oil (95% methyl caprylate)
Isopar® H (Exxon)	Isoparaffinic mixture, distillation range 182 - 192 °C
Shellisol® T (Shell)	Isoparaffinic mixture, C <sub>11</sub> - C <sub>13</sub>
Solvesso® 200 (Exxon)	Aromatic mixture, distillation range 231 - 287 °C
Attagel® 50 (Engelhard Corp.)	hydrated aluminium magnesium silicate

**Example A**

**Example B**

Ingredient	amount (g)	Ingredient	amount (g)
<u>Compound IB</u>	100	<u>Compound IB</u>	100
Rhodocal 70/B	50	Rhodocal 70/B	50
Ukanil 2507	50	Ukanil 2507	50
Plurafac LF700	480	Plurafac LF700	480
Witconol 1095	320	Isopar H	320

**Example C**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	50
Plurafac LF700	400
Shellsol T	400

**Example D**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	50
Plurafac LF700	480
Solvesso 200	320

Physico-chemical Properties

Property	Example			
	A	B	C	D
Blind formulation	clear	clear	clear	clear
Spray dilution of 0.5 ml formulation in graduated cylinder with 100 ml tap water, self-dispersion checked, followed by 30 inversions.	Good	good	good	good
Spray dilution 24 h, re-dispersibility of a.i. particles	good	good	good	good
Storage of formulation for 2 weeks at 54°C, visual evaluation of particles under microscope	no particle growth	no particle growth	no particle growth	no particle growth

**5 Example E**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	50
Attagel 50	60
Plurafac LF700	444
Solvesso 200	to 1 liter



Physico-chemical Properties

Property	Example E
Density	1.06 g/ml
Flash point	> 70°C
Spray dilution of 0.5 ml SC in graduated cylinder with 100 ml tap water, self-dispersion checked.	Good
Spray dilution 24 h, redispersibility of a.i. particles	good

**Example F****Example G**

Ingredient	amount (g)	Ingredient	amount (g)
<u>Compound IB</u>	100	<u>Compound IB</u>	100
Rhodocal 70/B	50	Rhodocal 70/B	50
Ukanil 2507	530	Ukanil 2507	50
Attagel 50	30	Attagel 50	30
Isopar H	to 1 liter	Plurafac LF700	480
		Isopar H	to 1 liter

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**Example H****Example I**

Ingredient	amount (g)	Ingredient	amount (g)
<u>Compound IB</u>	100	<u>Compound IB</u>	100
Rhodocal 70/B	50	Rhodocal 70/B	50
Ukanil 2507	50	Ukanil 2507	50
Attagel 50	30	Attagel 50	30
SCS4774	480	Emulgin CO3522	480
Isopar H	to 1 liter	Isopar H	to 1 liter

**Example J**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	530
Attagel 50	30
Witconol 1095	to 1 liter

**Example K**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	530
Attagel 50	30
Ethyl Diglyme	to 1 liter

**Example L**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	530
Attagel 50	30
Ethyl lactat	to 1 liter

**Example M**

Ingredient	amount (g)
<u>Compound IB</u>	100
Rhodocal 70/B	50
Ukanil 2507	530
Attagel 50	30
2-Ethylhexyl lactat	to 1 liter

What is claimed is:

1. A non-aqueous, stable suspension concentrate (SC) which comprises

(a) 50 to 400 g/L of one or more crop protection active compounds;

(b) 50 to 700 g/L of one or more adjuvants;

(c) 75 to 500 g/L of one or more organic solvents;

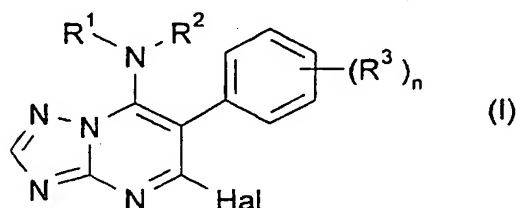
at least one dispersant selected from the groups (d) and (e)

(d) up to 150 g/L of one or more non-ionic dispersants,

(e) 5 to 150 g/L of one or more anionic dispersants; and optionally,

(f) up to 100 g/L of one or more thickeners.

2. A SC in accordance with claim 1 wherein said crop protection active compounds (a) comprise at least one triazolopyrimidine of formula I



in which

R<sup>1</sup> and R<sup>2</sup> each independently represent hydrogen or an optionally substituted alkyl, alkenyl, alkynyl, alkadienyl, haloalkyl, aryl, heteroaryl, cycloalkyl, bicycloalkyl or heterocyclyl group, or

R<sup>1</sup> and R<sup>2</sup> together with the adjacent nitrogen atom represent an optionally substituted heterocyclic ring,

R<sup>3</sup> represents a halogen atom or an alkyl or alkoxy group,

n represents an integer from 0 to 5, and

Hal represents a halogen atom.

3. A SC in accordance with claim 1 wherein said adjuvant (b) comprises a compound having 2 to 20 C<sub>2-6</sub> alkoxy groups.
4. A SC in accordance with claim 3 wherein said adjuvant (b) is a mixed ethoxylate/propoxylate.
5. A SC in accordance with claim 3 wherein said adjuvant (b) is selected from the group consisting of polyalkoxylated alcohols, amines and triglycerides.
6. A SC in accordance with claim 5 wherein said adjuvant (b) comprises two or more different polyalkoxylated derivatives, one of which is a polyalkoxylated triglyceride.
7. A SC in accordance with claim 1 wherein the ratio of said crop protection active compounds (a) to said adjuvant (b) is between 1:100 and 1:1.
8. A SC in accordance with claim 7 wherein said ratio is between 1:10 and 1:1.
9. A SC in accordance with claim 1 wherein the solvent (c) is selected from the group consisting of aromatic hydrocarbons, aliphatic hydrocarbons, alkyl lactates, glycols and plant oil esters, and mixtures thereof.
10. A SC in accordance with claim 1 wherein the non-ionic dispersant (d) is a polyoxyethylene fatty acid.
11. A SC in accordance with claim 1 wherein the anionic dispersant (e) is an alkali or alkaline earth metal sulfonate.

12. A SC in accordance with claim 1 wherein the thickener (f) is an organo clay.

13. A process for the preparation of a SC as claimed in claim 1, which comprises

- (a) air-milling of 50 to 400 g/L of one or more crop protection active compounds (a) optionally in the presence of one or more anionic dispersant (e), and/or a milling aid, and
- (b) mixing all the components (a) to (c), (d) and/or (e) and optionally (f) in a dissolver.

14. A method for the control of pests at a locus which comprises diluting a SC as claimed in claim 1 with water and treating the locus with a pesticidally effective amount of the diluted formulation.

## INTERNATIONAL SEARCH REPORT

onal Application No

PCT/US 99/22046

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A01N25/04 A01N43/90 //(A01N43/90,25:04)

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 789 999 A (HOECHST SCHERING AGREVO GMBH) 20 August 1997 (1997-08-20) page 2, line 18 - line 26 page 2, line 46 -page 3, line 36	1,3,5, 7-9,11, 12,14
X	EP 0 071 792 A (BASF AG) 16 February 1983 (1983-02-16) cited in the application page 18, line 26 - line 34 page 20, line 8 - line 13	1-3,5, 7-11,14
X	GB 2 083 360 A (DOW CHEMICAL CO) 24 March 1982 (1982-03-24) page 3, line 62 -page 4, line 21 page 4; examples 3,4	1,3-5, 7-12,14
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☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

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"P" document published prior to the international filing date but later than the priority date claimed

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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"8" document member of the same patent family

Date of the actual completion of the international search

2 February 2000

Date of mailing of the international search report

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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/22046

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 149 459 A (SIPCAM SPA) 24 July 1985 (1985-07-24)  page 1, line 12 - line 20 page 2, line 19 - page 3, line 20 page 5 - page 6; example 1 ---	1,3,5, 7-9,11, 12,14
X	EP 0 645 083 A (AMERICAN CYANAMID CO) 29 March 1995 (1995-03-29) page 2, line 37 - line 44 page 3, line 7 - line 29 page 3, line 44 - line 56 ---	1,3-5, 7-12,14
X	EP 0 313 317 A (ISHIHARA MINING & CHEMICAL CO) 26 April 1989 (1989-04-26) page 3, line 5 - line 19 page 4, line 1 - line 25 ---	1,5,7,9, 10,12,14
X	EP 0 456 198 A (HODOGAYA CHEMICAL CO LTD ;TOHO CHEM IND CO LTD (JP)) 13 November 1991 (1991-11-13) page 3, line 8 - line 32 page 3; example 1 ---	1,3-9, 11,13,14
X	EP 0 103 171 A (NISSAN CHEMICAL IND LTD) 21 March 1984 (1984-03-21) page 1, line 1 - line 7 page 3, line 13 - line 26 page 6, line 10 - line 12 page 7, line 5 - line 17 page 7, line 22 - line 24 ---	1,13,14
E	EP 0 943 241 A (AMERICAN CYANAMID CO) 22 September 1999 (1999-09-22) page 3, line 27 - line 57 page 10, line 1 - line 20 -----	1-3,5, 7-11,14

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/22046

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0789999	A	20-08-1997	DE 19605786 A	21-08-1997
			CZ 9700467 A	17-09-1997
			HU 9700446 A	28-10-1997
			PL 318490 A	18-08-1997
EP 0071792	A	16-02-1983	DE 3130633 A	17-02-1983
			AT 11539 T	15-02-1985
			AU 553663 B	24-07-1986
			AU 8665982 A	10-02-1983
			CA 1180329 A	01-01-1985
			CS 226748 B	16-04-1984
			DD 202093 A	31-08-1983
			DK 341682 A,B,	02-02-1983
			GR 76193 A	03-08-1984
			HU 188325 B	28-04-1986
			IE 53269 B	28-09-1988
			JP 1634879 C	20-01-1992
			JP 2061955 B	21-12-1990
			JP 58043974 A	14-03-1983
			US 4567263 A	28-01-1986
			ZA 8205498 A	27-07-1983
GB 2083360	A	24-03-1982	NONE	
EP 0149459	A	24-07-1985	IT 1173066 B	18-06-1987
			ZA 8500447 A	25-09-1985
EP 0645083	A	29-03-1995	US 5405825 A	11-04-1995
			AT 176382 T	15-02-1999
			CZ 9402356 A	12-04-1995
			DE 69416369 D	18-03-1999
			DE 69416369 T	10-06-1999
			ES 2127324 T	16-04-1999
			GR 3030122 T	30-07-1999
			HU 68866 A,B	28-08-1995
			SK 116594 A	12-04-1995
			US 5707928 A	13-01-1998
EP 0313317	A	26-04-1989	JP 1110605 A	27-04-1989
			JP 2569342 B	08-01-1997
			CN 1032729 A,B	10-05-1989
			DE 3870396 A	27-05-1992
			GR 3005151 T	24-05-1993
			HU 49446 A,B	30-10-1989
			KR 9306675 B	22-07-1993
			PL 275421 A	10-07-1989
			US H750 H	06-03-1990
EP 0456198	A	13-11-1991	JP 4018002 A	22-01-1992
			JP 2945076 B	06-09-1999
			JP 4021611 A	24-01-1992
			CN 1056216 A	20-11-1991
EP 0103171	A	21-03-1984	JP 1734948 C	17-02-1993
			JP 4017923 B	26-03-1992
			JP 59029604 A	16-02-1984
			AU 554234 B	14-08-1986
			AU 1736083 A	16-02-1984



## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/22046

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0103171 A		BR 8304356 A	20-03-1984
		CA 1211294 A	16-09-1986
EP 0943241 A	22-09-1999	JP 11322517 A	24-11-1999